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**UNITED STATES DISTRICT COURT
NORTHERN DISTRICT OF CALIFORNIA
SAN FRANCISCO DIVISION**

Louise Porras, an individual and on behalf of the
estate of Edward Porras,

Plaintiff,

v.

GlaxoSmithKline, LLC; Boehringer Ingelheim
Pharmaceuticals, Inc.; Boehringer Ingelheim USA
Corporation; Pfizer, Inc.; Sanofi US Services Inc.;
Sanofi-Aventis U.S. LLC; Patheon Manufacturing
Services, LLC; Walgreens Co.; Rite Aid
Corporation; and DOES 1 through 100 inclusive,

Defendants.

Case No.

**NOTICE OF REMOVAL OF ACTION
PURSUANT TO 28 U.S.C. §§ 1332, 1441,
and 1446**

DEMAND FOR JURY TRIAL

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Pursuant to 28 U.S.C. §§ 1332, 1441, and 1446, Defendants GlaxoSmithKline, LLC (“GSK”), Pfizer Inc. (“Pfizer”), Boehringer Ingelheim Pharmaceuticals, Inc. and Boehringer Ingelheim USA Corporation (together, “BI”), Sanofi US Services Inc. and Sanofi-Aventis U.S. LLC (together, “Sanofi”), Walgreen Co. (“Walgreens”), and Rite Aid Corporation (“Rite Aid”) (collectively, “Removing Defendants”) hereby give notice of removal of this action, *Porrás v. GlaxoSmithKline, LLC et al.*, Case No. CGC22603510, from the Superior Court of the State of California in and for the County of San Francisco to the United States District Court for the Northern District of California.

INTRODUCTION

1. This action is one of many individual lawsuits filed against manufacturers, distributors, and retailers of Zantac (ranitidine) relating to cancers allegedly caused by the drug. *See* Compl., attached as **Exhibit A**.¹ On February 6, 2020, the Judicial Panel on Multidistrict Litigation (“JPML”) created a Multidistrict Litigation (“MDL”) in the Southern District of Florida before Judge Robin Rosenberg for pretrial coordination of cases like this one “in which plaintiffs allege that they developed cancer as a result of NDMA formed from Zantac.” *In re Zantac (Ranitidine) Prods. Liab. Litig.*, 437 F. Supp. 3d 1368, 1369 (J.P.M.L. 2020). The JPML found that centralizing these cases for pretrial purposes “will eliminate duplicative discovery; prevent inconsistent pretrial rulings . . . and conserve the resources of the parties, their counsel, and the judiciary.” *Id.* To date, thousands of actions have been filed in or transferred to the Zantac MDL.

2. In this case, the only non-diverse defendant, Patheon Manufacturing Services, LLC (“Patheon”), is fraudulently joined. Prior to filing this lawsuit, Plaintiff asserted in a verified statement submitted in the Zantac MDL that [REDACTED]

[REDACTED] *See* Pl.’s Census Plus Form at [REDACTED], attached hereto as **Exhibit B**. But Patheon never manufactured, packaged, distributed, or sold [REDACTED]

[REDACTED],

¹ The Complaint alleges that the decedent consumed ranitidine products, and, as a result, developed colorectal cancer. *See* Compl. ¶ 9. Plaintiff asserts eight claims against the Removing Defendants: (1) strict liability – failure to warn; (2) strict liability – manufacturing defect; (3) negligence – failure to warn; (4) negligent product design; (5) negligent manufacturing; (6) negligent misrepresentation; (7) wrongful death; and (8) survival.

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Patheon therefore could not have manufactured or distributed

3. As such, Plaintiff cannot state a cause of action against Patheon. Patheon's citizenship must be disregarded, and federal jurisdiction exists because: (1) there is complete diversity of citizenship between Plaintiff and all properly joined defendants; and (2) the amount in controversy exceeds the jurisdictional threshold under 28 U.S.C. § 1332(a).

VENUE AND JURISDICTION

4. Venue is proper in this Court pursuant to 28 U.S.C. §§ 84(a), 1391, 1441(a), and 1446(a) because the Superior Court of the State of California in and for the County of San Francisco, where the Complaint was filed, is a state court within the Northern District of California.

5. This Court has subject matter jurisdiction under 28 U.S.C. § 1332(a) because: (1) there is complete diversity between Plaintiff and all properly joined defendants; (2) the amount in controversy exceeds \$75,000, exclusive of interest and costs; and (3) all other requirements for removal have been satisfied.

BASIS FOR REMOVAL**I. There Is Complete Diversity Between Plaintiff and All Properly Joined Defendants.**

6. In a representative lawsuit such as this one, "the legal representative of the estate of a decedent shall be deemed to be a citizen only of the same State as the decedent." 28 U.S.C. § 1332(c)(2). Plaintiff alleges that she is the "surviving heir[], successor[] in interest, and/or personal representative[] of Decedent[]," who in turn "was a citizen and resident of the state of California." Compl. ¶¶ 9, 22, 282, 288. Plaintiff is, therefore, a citizen of California.

7. For purposes of diversity jurisdiction, a corporation is "a citizen of every State and foreign state by which it has been incorporated and of the State or foreign state where it has its principal place of business." 28 U.S.C. § 1332(c)(1). A limited liability company "is a citizen of every state of which its owners/members are citizens." *Johnson v. Columbia Props. Anchorage, LP*, 437 F.3d 894, 899 (9th Cir. 2006).

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1 8. Defendant Boehringer Ingelheim Pharmaceuticals, Inc. is a corporation organized
2 under the laws of Delaware with its principal place of business in Ridgefield, Connecticut. Compl.
3 ¶ 27. Boehringer Ingelheim Pharmaceuticals, Inc. is, therefore, a citizen of Delaware and
4 Connecticut.

5 9. Defendant Boehringer Ingelheim USA Corporation is a corporation organized under
6 the laws of Delaware with its principal place of business in Ridgefield, Connecticut. *Id.* ¶ 28.
7 Boehringer Ingelheim USA Corporation is, therefore, a citizen of Delaware and Connecticut.

8 10. Defendant GlaxoSmithKline LLC is a limited liability company organized under the
9 laws of Delaware. *Id.* ¶ 23. GlaxoSmithKline LLC's sole member is GlaxoSmithKline Holdings
10 (America) Inc., a corporation organized under the laws of Delaware with its principal place of
11 business in Wilmington, Delaware. *See* Attachment to Corp. Disclosure Statement, *In re Zantac*
12 *(Ranitidine) Prods. Liab. Litig*, MDL No. 2924, ECF No. 43-1. GlaxoSmithKline LLC is, therefore,
13 a citizen of Delaware.

14 11. Defendant Pfizer Inc. is a corporation organized under the laws of Delaware with its
15 principal place of business in New York, New York. Compl. ¶ 26. Pfizer Inc. is, therefore, a citizen
16 of Delaware and New York.

17 12. Defendant Sanofi US Services Inc. is a corporation organized under the laws of
18 Delaware with its principal place of business in Bridgewater, New Jersey. *Id.* ¶ 30. Sanofi US
19 Services Inc. is, therefore, a citizen of Delaware and New Jersey.

20 13. Defendant Sanofi-Aventis U.S. LLC is a limited liability company organized under
21 the laws of Delaware. *Id.* ¶ 31. The sole member of Sanofi-Aventis U.S. LLC is Sanofi US Services
22 Inc., a Delaware corporation with its principal place of business in Bridgewater, New Jersey. *Id.*
23 ¶ 30. Sanofi-Aventis U.S. LLC is, therefore, a citizen of Delaware and New Jersey.

24 14. Defendant Walgreen Co. is a corporation organized under the laws of Illinois with its
25 principal place of business in Deerfield, Illinois. *Id.* ¶ 34. Walgreen Co. is, therefore, a citizen of
26 Illinois.

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1 15. Defendant Rite Aid Corporation is a corporation organized under the laws of
2 Delaware with its principal place of business in Camp Hill, Pennsylvania. *Id.* ¶ 35. Rite Aid
3 Corporation is, therefore, a citizen of Delaware and Pennsylvania.

4 16. Defendants “Does 1 through 100” are disregarded when assessing the diversity of
5 citizenship of the parties for removal. *See* 28 U.S.C. § 1441(b)(1) (“In determining whether a civil
6 action is removable on the basis of the jurisdiction under section 1332(a) of this title, the citizenship
7 of defendants sued under fictitious names shall be disregarded.”); *see also Soliman v. Philip Morris*
8 *Inc.*, 311 F.3d 966, 971 (9th Cir. 2002) (“The citizenship of fictitious defendants is disregarded for
9 removal purposes and becomes relevant only if and when the plaintiff seeks leave to substitute a
10 named defendant.”).

11 **II. Patheon Is Fraudulently Joined.**

12 17. Patheon is alleged to be a citizen of Delaware, New York, California, Massachusetts,
13 Wisconsin, and Pennsylvania through a complicated corporate structure. *See* Compl. ¶ 33. But
14 because Patheon is fraudulently joined in this action, its citizenship should be disregarded.

15 18. A defendant is fraudulently joined and its citizenship can be ignored for establishing
16 diversity jurisdiction, where no viable cause of action has been stated against it. *See Morris v.*
17 *Princess Cruises, Inc.*, 236 F.3d 1061, 1067 (9th Cir. 2001) (a defendant is fraudulently joined “[i]f
18 the plaintiff fails to state a cause of action against a resident defendant, and the failure is obvious
19 according to the settled rules of the state”) (quoting *McCabe v. Gen. Foods Corp.*, 811 F.2d 1336,
20 1339 (9th Cir. 1987)); *Ritchey v. Upjohn Drug Co.*, 139 F.3d 1313, 1318 (9th Cir. 1998) (same).

21 19. Under California law, a plaintiff asserting a personal injury product liability claim
22 must demonstrate that “the defendant’s product, or some instrumentality under the defendant’s
23 control, caused his or her injury.” *DiCola v. White Bros. Performance Prods., Inc.*, 69 Cal. Rptr.
24 3d 888, 897 (Ct. App. 2008) (collecting cases); *see also Garcia v. Joseph Vince Co.*, 148 Cal. Rptr.
25 843, 846 (Ct. App. 1978) (“[W]hether negligence, . . . strict liability in tort, or other grounds, it is
26 obvious that to hold a producer, manufacturer, or seller liable for injury caused by a particular
27 product, there must first be proof that the defendant produced, manufactured, sold, or was in some
28 way responsible for the product.” (quoting 51 A.L.R.3d 1334, 1349)); *Soule v. Gen. Motors Corp.*,

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1 882 P.2d 298, 312 (Cal. 1994) (“A manufacturer is liable only when a defect in its product was a
 2 legal cause of injury.” (citation omitted)); *Sanchez v. Stryker Corp.*, No. 2:10-CV-08832-ODW,
 3 2012 WL 1570569, at *6 & n.9 (C.D. Cal. May 2, 2012) (“These five causes of action [including
 4 negligence, strict liability, and negligent misrepresentation] have one central question—whether the
 5 specific Stryker device . . . caused Plaintiff’s injury.” (collecting cases)); *Timmons v. Linvatec Corp.*,
 6 263 F.R.D. 582, 585 (C.D. Cal. 2010) (dismissing negligence, strict liability, negligent
 7 misrepresentation, and consortium claims against pharmaceutical manufacturer with prejudice, in
 8 part because it was undisputed that the defendant had not manufactured or sold the drug at issue).
 9 In other words, Plaintiff must establish that a particular defendant’s product caused her injury in
 10 order to succeed on any of her claims against that defendant.

11 20. Fraudulent joinder claims can be resolved by “piercing the pleadings” and considering
 12 outside evidence. *See Morris*, 236 F.3d at 1068 (quoting *Cavallini v. State Farm Mutual Auto Ins.*
 13 *Co.*, 44 F.3d 256, 263 (5th Cir. 1995) (“[F]raudulent joinder claims may be resolved by ‘piercing
 14 the pleadings’ and considering summary judgment-type evidence such as affidavits and deposition
 15 testimony.”)); *BSD, Inc. v. Equilon Enters., LLC*, No. C 10-05223 SBA, 2011 WL 1295984, at *3
 16 (N.D. Cal. Apr. 1, 2011) (same).

17 21. Applying these principles, courts routinely find that non-diverse defendants in product
 18 liability actions are fraudulently joined where those defendants did not actually manufacture or sell
 19 the product that allegedly caused the plaintiff’s injury. *See, e.g., Gallagher v. Boehringer Ingelheim*
 20 *Pharms., Inc.*, No. 22-CV-10216 (LJL), 2023 WL 402191, at *5–6, 10 (S.D.N.Y. Jan. 25, 2023)
 21 (denying remand after finding that non-diverse defendant was fraudulently joined because that
 22 defendant “was no longer in control of Zantac when [the drug] caused Plaintiff’s injuries”); *Vieira*
 23 *v. Mentor Worldwide, LLC*, 392 F. Supp. 3d 1117, 1127–28 (C.D. Cal. 2019) (finding non-diverse
 24 defendant fraudulently joined because it “did not manufacture . . . and was not involved in the
 25 development of the MemoryGel Implant,” and therefore there was “no possibility that Plaintiff
 26 could recover under a theory of product liability against [it]”), *aff’d*, 845 F. App’x 503 (9th Cir.
 27 2021); *Reith v. Teva Pharms. USA, Inc.*, No. CV 18-3987, 2019 WL 1382624, at *3-5 (E.D. Pa.
 28 Mar. 27, 2019) (dismissing as fraudulently joined non-diverse defendants that were “not in the chain

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1 of distribution as the manufacturer or seller of the plaintiffs' IUDs" as required in order to state a
 2 claim against them under state law, and denying remand).

3 22. Here too, Patheon is fraudulently joined, because Patheon did not manufacture,
 4 package, distribute, or sell [REDACTED]

5 23. [REDACTED]
 6 [REDACTED].

7 24. [REDACTED]
 8 [REDACTED] [REDACTED]
 9 [REDACTED]
 10 [REDACTED]
 11 [REDACTED]
 12 [REDACTED].²

13 25. While the Complaint fails to specify the ranitidine products that the decedent took,
 14 prior to filing this case, Plaintiff asserted that the [REDACTED] in the
 15 "Census Plus Form" that she verified and submitted in the Zantac MDL. In response to questions
 16 seeking information about [REDACTED] Plaintiff answered that [REDACTED]
 17 [REDACTED] See Exhibit B at
 18 [REDACTED] (emphasis in original)). When asked to [REDACTED]
 19 [REDACTED]
 20 [REDACTED] Plaintiff certified
 21 that "under penalty of perjury" her answers on the Census Plus Form were "true and correct." *Id.*
 22 at 6.

23 26. Because Plaintiff has asserted that [REDACTED], Ex. B
 24 at [REDACTED], no Patheon product could possibly have caused the decedent's alleged injuries. As such,
 25 Plaintiff cannot prevail on any of her claims against Patheon. Thus, Patheon is fraudulently joined
 26

27 ² The Complaint also asserts claims against Walgreens and Rite Aid, alleging that these retailers
 28 "operat[ed] [] pharmac[ies] which dispense[] Ranitidine-Containing Drugs" in California. Compl.
 ¶¶ 34, 35.

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1 and its citizenship can be ignored for establishing diversity jurisdiction.³ *See Morris*, 236 F.3d at
2 1067. When Patheon's citizenship is ignored, Plaintiff is diverse from all properly joined
3 defendants. *See* 28 U.S.C. §§ 1332, 1441.

III. The Amount-in-Controversy Requirement Is Satisfied

5 27. Plaintiff's claims satisfy the amount in controversy requirement set forth in 28 U.S.C.
6 § 1332(a).

7 28. California Code of Civil Procedure § 425.10(b) prevents a plaintiff from stating a
8 specific damage claim in a personal injury action.

9 29. "[A] defendant's notice of removal need include only a plausible allegation that the
10 amount in controversy exceeds the jurisdictional threshold." *Dart Cherokee Basin Operating Co.,*
11 *LLC v. Owens*, 135 S. Ct. 547, 554 (2014). Where a complaint does not set forth a specific amount
12 of damages, it may nevertheless be "facially apparent" that the amount in controversy exceeds the
13 jurisdictional minimum. *See Mobasser v. Travelers Cas. Ins. Co. of Am.*, No. 13-CV-02567-DMG-
14 CW, 2013 WL 12142942, at *2 (C.D. Cal. June 13, 2013) (quoting *Singer v. State Farm Mut. Auto.*
15 *Ins. Co.*, 116 F.3d 373, 377 (9th Cir. 1997)). "[T]he defendant's amount-in-controversy allegation
16 should be accepted when not contested by the plaintiff or questioned by the court," and "[e]vidence
17 establishing the amount is required by § 1446(c)(2)(B) only when the plaintiff contests, or the court
18 questions, the defendant's allegation." *Dart Cherokee*, 135 S. Ct. at 553–54.

19 30. Here, Plaintiff seeks several categories of damages, including actual or compensatory
20 damages, exemplary damages, punitive damages, and attorneys' fees. *See* Compl., Prayer for Relief
21 ¶ 292(a)-(e). The Complaint includes eight causes of action and alleges that the decedent's use of
22 Zantac caused him to suffer "significant harm, conscious pain and suffering, physical injury and
23 bodily impairment including, but not limited to cancer, other permanent physical deficits, permanent
24

25 ³ While not material to the jurisdictional analysis, based on the decedent's [REDACTED]
26 [REDACTED] Pfizer, BI, and Sanofi are also improper defendants. Pfizer, BI, and Sanofi
only manufactured, marketed, and/or sold [REDACTED]

27 [REDACTED] However, because
28 Pfizer, BI, and Sanofi are not forum defendants and are completely diverse from Plaintiff, this Court
need not determine whether they are fraudulently joined for purposes of removal.

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1 bodily impairment” and death, as well as “mental anguish, [] loss of enjoyment of life . . . [and]
2 economic losses.” *Id.* ¶¶ 12, 14, 15, 21. It further asserts that these injuries “required
3 hospitalizations, in-patient surgeries, medication treatments, and other therapies,” which resulted in
4 “loss of earning and medical expenses.” *Id.* ¶¶ 12, 21. And it alleges that Plaintiff suffered
5 “depriv[ation] of . . . companionship.” *See id.* ¶ 285.

6 31. Courts regularly find that the amount-in-controversy requirement is apparent from the
7 face of the complaint where, as here, a plaintiff alleges serious bodily injury and death. *See, e.g.,*
8 *Campbell v. Bridgestone/Firestone, Inc.*, No. 05-CV-01499-FVS-DLB, 2006 WL 707291, at *2–3
9 (E.D. Cal. Mar. 17, 2006) (amount in controversy met where complaint asserted strict products
10 liability, negligence, and breach of warranty claims against multiple defendants and requested
11 compensatory damages, hospital and medical expenses, general damages, and loss of earnings);
12 *Geographic Expeditions, Inc. v. Estate of Lhotka*, 599 F.3d 1102, 1107–08 (9th Cir. 2010) (amount
13 in controversy met where complaint requested damages for, among other things, wrongful death,
14 loss of consortium, and negligence, as well as funeral, medical and burial expenses); *Mullaney v.*
15 *Endogastric Sols. Inc.*, No. 11-62056-CIV, 2011 WL 4975904, at *2 (S.D. Fla. Oct. 19, 2011)
16 (inferring amount-in-controversy requirement was met where plaintiff alleged that he “underwent
17 ‘surgical intervention that required additional life-saving medical treatment’ and suffered ‘serious,
18 permanent and disabling injuries’”); *In re Rezulin Prods. Liab. Litig.*, 133 F. Supp. 2d 272, 296–97
19 (S.D.N.Y. 2001) (concluding jurisdictional amount “obviously” met where plaintiff sought to
20 recover punitive and compensatory damages, including lost wages and medical expenses for serious
21 and life-threatening conditions).

22 32. Indeed, in another Zantac case in which the plaintiff claimed cancer as an injury, a
23 federal court in this Circuit denied a motion to remand where the amount in controversy was not
24 alleged, finding that the requirement was satisfied on the face of the complaint by the nature of the
25 injury. *See Brooks v. Sanofi S.A.*, No. 20-CV-565, 2020 WL 1847682 (D. Nev. Apr. 13, 2020).

26 33. Moreover, California courts have awarded compensatory and punitive damages in
27 excess of \$75,000 in products liability cases where similar injuries were alleged. *See, e.g., Karlsson*
28

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1 *v. Ford Motor Co.*, 45 Cal. Rptr. 3d 265, 268 (Ct. App. 2006); *Jones v. John Crane, Inc.*, 35 Cal.
2 Rptr. 3d 144, 148–49 (Ct. App. 2005).

3 34. Finally, in the hundreds of personal injury cases pending in the Zantac MDL, each
4 plaintiff either expressly claims damages in excess of \$75,000 or has impliedly done so by filing a
5 lawsuit in federal court and invoking federal diversity jurisdiction. Numerous plaintiffs in these
6 cases allege that they have been diagnosed with colorectal cancer, the same type of cancer alleged
7 in the Complaint. *See, e.g., Roman-Padin v. GlaxoSmithKline LLC et al.*, No. 9:21-cv-82084 (S.D.
8 Fla.); *Howell v. Sanofi-Aventis U.S. LLC et al.*, No. 9:21-cv-80101 (S.D. Fla.).

9 35. Although the Removing Defendants deny that Plaintiff is entitled to any damages, the
10 allegations in Plaintiff’s Complaint demonstrate that the amount in controversy exceeds \$75,000,
11 exclusive of interest and costs.

12 **IV. Procedural Requirements of Removal Are Satisfied**

13 36. This Notice of Removal is timely filed pursuant to 28 U.S.C. § 1446(b).

14 37. Boehringer Ingelheim Pharmaceuticals, Inc. and Boehringer Ingelheim USA
15 Corporation were served with the Complaint on January 24, 2023.

16 38. GlaxoSmithKline, LLC, Pfizer Inc., Sanofi US Services Inc., Sanofi-Aventis U.S.
17 LLC, Walgreen Co., and Rite Aid Corporation were served with the Complaint on January 25, 2023.

18 39. Removal pursuant to 28 U.S.C. § 1441(a) requires that “all defendants who have been
19 properly joined and served must join in or consent to the removal of the action.” 28 U.S.C.
20 § 1446(b)(2)(A).

21 40. All of the Removing Defendants consent to this removal.

22 41. Patheon is fraudulently joined and therefore is not required to join in the removal. *See*
23 *United Computer Sys. Inc. v. AT&T Corp.*, 298 F.3d 756, 762 (9th Cir. 2002).

24 42. The unidentified defendants Does 1–100 are not required to consent to removal. *See*
25 *Hafiz v. Greenpoint Mortg. Funding*, 409 F. App’x 70, 72 (9th Cir. 2010) (nominal parties are not
26 required to consent to removal).

27 43. The Removing Defendants are providing Plaintiff with written notice of the filing of
28 this Notice of Removal as required by 28 U.S.C. § 1446(d).

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